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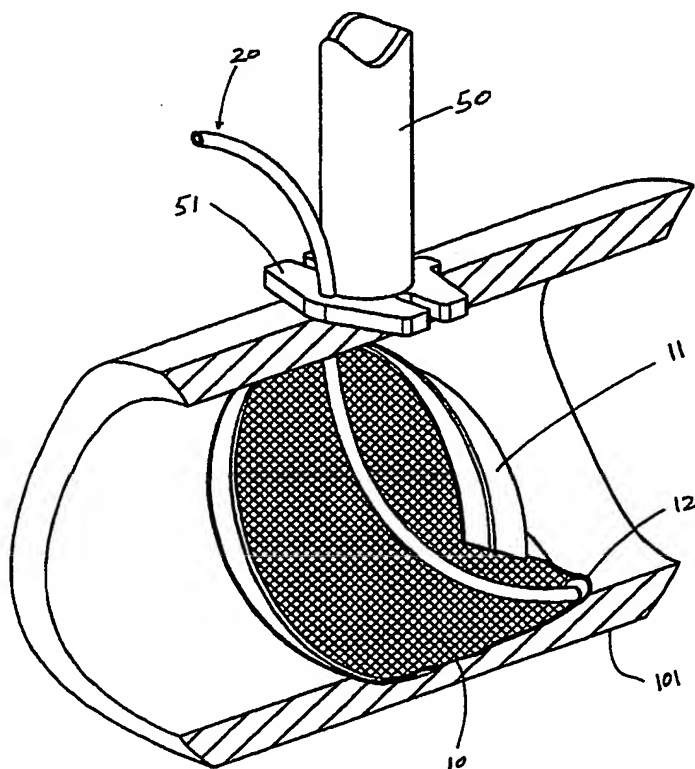
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(54) Title: ARTERIAL FILTER WITH ASPIRATION AND METHODS OF USE



(57) Abstract: A filter (10) comprising a porous material is arranged in a generally conical shape having an expandable base (11) and a vertex (12). A reservoir tip (15) is located at the vertex (12) for collecting filtered embolic material. The tip (15) communicates with a distal port (26) and lumen of an aspiration tube (20) for removing embolic material under vacuum. A vessel occluding device (60) comprising a non-permeable material having an expandable conical shape and communicating with an aspiration tube (20) is also disclosed. Methods of using the filter and occluding devices for protecting a patient from embolization during cardiovascular procedures are described.

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Arterial Filter with Aspiration and Methods of Use

Field of the Invention

5 The present invention relates generally to medical devices for protecting a patient from embolization during cardiovascular procedures. More particularly, the devices comprise a filter for temporary placement in a patient's vessel, and an aspiration tube communicating with a portion of the filter, *e.g.*, a reservoir tip at a vertex of the filter, for aspirating embolic material, including air and gas.

10 Background of the Invention

Aspiration catheters are frequently used during surgical or interventional procedures for removing thromboembolic material, *e.g.*, air, fluid, thrombi, calcium, atheromatous plaque, and/or tissue debris, from a patient's body cavity. During cardiovascular procedures, such as coronary artery bypass grafting surgery, ventricular
15 septal defect repair, heart valve repair or replacement, ventricular myomectomy, aortic aneurysm repair, or aortic thrombectomy, removal of thromboembolic material from a cardiac chamber and/or the aorta is important since distal embolization may result in ischemia or infarction of peripheral organs, particularly the brain.

Endovascular techniques have been used widely as an effective and reliable
20 alternative to surgical intervention in selected patients for treatment of vascular stenosis. It is well recognized that one of the complications associated with endovascular techniques is the dislodgment of embolic materials generated during manipulation of the vessel, thereby causing occlusion of the narrower vessels downstream and ischemia or infarct of the organ which the vessel supplies.

25 Several arterial and venous filters have been designed for entrapment of embolic debris generated during surgical or endovascular procedures. An aspiration catheter is often required for complete removal of embolic debris. Current aspiration catheters are designed to remove fluid and tissue debris in a body cavity. Removal of air, however, is difficult because the air bubbles tend to accumulate against the vessel wall at a position

difficult to reach. Thus, removal of embolic material is often not complete and patients remain at risk for air embolization.

Thus, there is a need for devices and methods which are capable of capturing and effectively removing embolic material within a patient's body tissue or cavity during surgical or endovascular procedures.

Summary of the Invention

The present invention provides an intravascular filter for temporary placement in a patient's vessel, such as an artery or vein. In a first embodiment, the filter comprises a porous material arranged in a generally conical shape having a base and a vertex. The base is expandable and collapsible. A reservoir tip is located at the vertex of the cone for collecting filtered embolic material. The tip also communicates with a distal end of an aspiration catheter. In certain embodiments, the reservoir tip comprises a nonporous material, *e.g.*, latex. In other embodiments, the distal end of the aspiration tube extends only to the outer edge of the filter, *e.g.*, the point of insertion into the vessel.

In another embodiment, the filter is mounted on a cannula, and the aspiration tube extends proximally within the cannula. The proximal end of the aspiration tube communicates with a vacuum pump.

In another embodiment, the filter is mounted on an insertion handle, and the aspiration tube extends proximally within the insertion handle. The aspiration tube includes an infusion port at the distal end for infusion of fluid, such as saline or heparin.

In another embodiment, the filter is mounted on a distal end of an obturator, and the aspiration tube extends proximally within the obturator. The distal end of the aspiration tube may further include a turbine that extends into the reservoir tip for removing large embolic particles.

The present invention further provides occluding devices for temporary placement in a patient's vessel, such as an artery or vein. In a first embodiment, the device comprises a nonporous material arranged in a conical shape having an expandable base and a vertex. A reservoir tip is located at the vertex of the cone for collecting filtered embolic material. The tip also communicates with a distal end of an aspiration catheter. In certain embodiments, the device is mounted on an insertion handle, having the aspiration tube

extending proximally within the handle. Alternatively, the device is mounted on a cannula or an obturator.

The invention also provides methods for protecting a patient from embolization using the filter or occluding devices described above. During cardiopulmonary bypass, for example, the filter or the occluding device is placed in a collapsed state and inserted into the patient's aorta. The filter or the occluding device is expanded to capture embolic material, including air, fluid, thrombi, calcium, atheromatous plaque, and/or tissue debris. The proximal end of the aspiration catheter is connected to a vacuum, and emboli are aspirated under negative pressure and removed from the aorta. The filter or the occluding device is then collapsed and removed from the aorta.

It will be understood that there are several advantages associated in using the devices and methods disclosed herein for preventing embolic complication during cardiovascular procedures. For example, (1) the filter is adapted for temporary placement in a patient's artery or vein; (2) the filter communicates with an aspiration tube for removal of embolic material; (3) the filter includes capability, such as a turbine or infusion port, for breakup of large embolic particles; (4) the occluding device provides aortic occlusion during cardiopulmonary bypass in addition to providing protection from distal embolization; (5) using the occluding device in place of an aortic clamp minimizes vascular injury caused by the clamp; and (6) the devices remove air and gas in addition to particles liable to cause distal embolization.

Brief Description of the Drawings

Fig. 1A depicts a filter with aspiration capabilities deployed in a vessel.

Fig. 1B depicts a lateral view of a filter having a reservoir tip communicating with an aspiration catheter.

Fig. 1C depicts a distal view of the filter of Fig. 1B.

Fig. 2A depicts a lateral view of another embodiment of the filter including a conical-shaped nonporous material.

Fig. 2B depicts a distal view of the filter of Fig. 2A.

Fig. 3A depicts another embodiment of the filter having an infusion port.

Fig. 3B depicts another embodiment of the filter having an angulated infusion port.

Fig. 4 depicts another embodiment of the filter having a turbine at a distal end of the aspiration catheter.

Fig. 5 depicts the filter of Fig. 1A passing through an introducer.

Fig. 6 depicts an occluder deployed in a vessel.

5 Fig. 7 depicts the filter of Fig. 1A mounted on a distal end of an insertion handle.

Fig. 8 depicts the filter of Fig. 1A deployed through an obturator.

Fig. 9 depicts the filter of Fig. 1A mounted on a distal end of a cannula.

Fig. 10 depicts a dam deployed in the aorta during cardiopulmonary bypass.

10 Detailed Description

The devices and methods disclosed herein are adapted for temporary placement in a patient's artery or vein for entrapment and removal of embolic debris. The devices are particularly useful in cardiovascular surgeries for preventing distal embolization to peripheral organs. An embodiment of the filter is depicted in Figs. 1A, 1B, and 1C. Filter
15 10 comprises a porous material arranged in a conical shape having expandable base 11 and vertex 12. Reservoir tip 15 is located at vertex 12 for collecting embolic material, and communicates with distal port 26 at distal end 22 and lumen 25 of aspiration tube 20. A proximal end (not shown) of the aspiration tube is adapted for attachment to a vacuum. In this embodiment, reservoir tip 15 comprises a nonporous material, *e.g.*, latex. In other
20 embodiments, the reservoir tip is constructed of the same material as the filter mesh.

The construction and use of expansion means and associated filter mesh have been thoroughly discussed in earlier applications including Barbut et al., U.S. Application Ser. No. 08/533,137, filed November 7, 1995, Barbut et al., U.S. Application Serial No. 08/580,223, filed December 28, 1995, Barbut et al., U.S. Application Serial No.
25 08/584,759, filed January 9, 1996, Barbut et al., U.S. Patent No. 5,769,816, filed April 30 1996, Barbut et al., U.S. Application Serial No. 08/645,762, filed May 14, 1996, and Barbut et al., U.S. Patent No. 5,662,671, and the contents of each of these prior applications are expressly incorporated herein by reference in their entirety.

Once appropriate physical characteristics are determined, suitable mesh can be
30 found among standard meshes known in the art. For example, polyester meshes may be used, such as meshes made by Saati Corporations and Tetko Inc. These are available in

sheet form and can be easily cut and formed into a desired shape. In a preferred embodiment, the mesh is sonic welded into a cone shape. Other meshes known in the art, which have the desired physical characteristics, are also suitable. For example, the filter may be constructed of a thin film material with laser cut holes. Anticoagulant, such as heparin and heparinoids, may be applied to the mesh to reduce the chances of blood clotting on the mesh. Anticoagulants other than heparinoids also may be used, e.g., ReoPro (Centocor). The anticoagulant may be painted or sprayed onto the mesh. A chemical dip comprising the anticoagulant also may be used. Other methods known in the art for applying chemicals to mesh may be used.

Another embodiment of the filter is depicted in Figs. 2A and 2B. Reservoir tip 15 comprises a nonporous material, e.g., latex, which extends into a portion of filter 10 and covers area 30. Area 30 is shaped like a sector of a circle having two radii and the included arc. In certain embodiments, latex may extend to cover distal end 22 of aspiration tube 20 as shown in Fig. 2B.

Fig. 3A depicts another embodiment of aspiration tube 20 having infusion port 40 at distal end 22. Infusion port 40 communicates with infusion lumen 41, which is included in the aspiration catheter. The infusion port and lumen are adapted for infusion of fluid, e.g., saline, or Ringer's lactate, for irrigating embolic material, and pharmaceutical agent, e.g., heparin or streptokinase to facilitate breakup of large thrombi dislodged within tip 15. Infusion port 40 may be angled relative to the axis of infusion lumen 41 as depicted in Fig. 3B.

In another embodiment, distal end 22 of aspiration tube 20 includes turbine 44 as shown in Fig. 4. The turbine can be activated proximally to disintegrate or break up large embolic material to facilitate removal into lumen 25.

In using the filter and aspiration device of Figs. 1A-1C, the filter can be introduced intravascularly directly through an open incision on the vessel. More preferably, the filter is placed in a collapsed state and inserted through lumen 55 of introducer 50 as depicted in Fig. 5. Introducer 50 is first introduced through an incision on vessel 100. Suture(s) may be placed on flange 51 to stabilize the introducer onto vessel 100. Filter 10 is then inserted into lumen 55 and deployed in the vessel. Aspiration tube 20 extends proximally within introducer 50 and is connected to a vacuum at its proximal end. The filter is expanded at its base, which covers the perimeter of the vessel wall. The filter captures and

the reservoir tip collects embolic material, e.g., air, fluid, thrombi, calcium, atheromatous plaque, and/or tissue debris, generated during the procedure. Emboli are aspirated into lumen 35 of aspiration tube 20 from the reservoir tip under negative pressure and removed. Emboli are also removed after the filter is collapsed and retrieved from
5 introducer 50. In this way, removal of embolic material liable to cause distal embolization during cardiovascular procedures is maximized.

Fig. 6 depicts vessel occluding device 60 deployed in aorta 101. The occluder comprises a non-permeable or semi-permeable material arranged in a generally conical shape having expandable base 11 and vertex 12. Reservoir tip 15 is located at vertex 12
10 for collecting embolic material. Distal port 26 and lumen 25 of aspiration tube 20 communicates with tip 15. The device is most useful in cardiovascular surgeries requiring cardiopulmonary bypass during isolation of coronary arteries from the peripheral vascular circulation. Using occluding device 60 in place of an aortic clamp, injury to the aortic wall, such as dissection and hematoma, can be minimized.

15 In use, during cardiopulmonary bypass, for example, occluder 60 is placed in a collapsed state and introduced into aorta 101 upstream of the placement of aortic cannula 110. Occluder 60 is then expanded to occlude the aortic lumen. Oxygenated blood is infused, generally at approximately 3 to 4 liters/minute, through aortic cannula 110 downstream in aorta 101 to perfuse peripheral organs. Embolic material generated during
20 cardiovascular surgery is collected in reservoir tip 15 of occluder 60. Since the heart is arrested during the procedure, pressure in the aorta distal to occluder 60 is higher than the pressure proximal to the occluder. As a result, blood flows from the high pressure side to the low pressure side, and in absence of aspiration, this reverse flow prevents the particles of emboli from reaching the filter, instead causing embolic particles to build up in the
25 coronary arteries. With aspiration, however, blood and the embolic material are removed under vacuum through aspiration tube 20.

Occluder 60 is activated to provide aspiration at cross-clamp removal such that a slightly lower pressure would exist in the ascending aorta proximal to occluder 60 and distal to the region from where the cross-clamp was removed. Blood flows from the high
30 pressure downstream portion of the aorta that is perfused by arterial return cannula 110 to upstream of occluder 60 because (1) small gaps are present between the vessel wall and the isolating filter/dam 60, and/or (2) the isolating filter/dam 60 is semi-permeable. The

permeable portion is designed to enhance flow into the ascending aorta proximal occluder 60, causing particles to be swept into the aspiration zone for removal. In another method, occluder 60 is non-permeable and cuffed with a balloon, thereby removing the need for a cross-clamp.

5 In another method, the proximal end of the aspiration tube is attached to a vacuum and an extracorporeal filter. In this way, blood or fluid is removed from the proximal aorta at a rate of 0.5 to 1 liter/min., is filtered free of debris, and is returned to the distal aorta through aortic cannula 110 at a rate of 3 to 4 liters/min., thereby minimizing blood loss.

10 Filter 10 can be mounted on a distal end of insertion handle 65 and inserted into access lumen 66 of cannula 110 as depicted in Fig. 7. In use, cannula 110 and access port 67 are inserted through an incision on aorta 101. Suture(s) may be placed on flange 68 to secure the cannula onto aorta 101. Filter 10 in a collapsed state is inserted through access lumen 66 and deployed in the aorta. Aspiration tube 20 extends proximally within
15 insertion handle 65 and is connected through proximal end 69 of the handle to pressure monitor 70, which is then connected to vacuum 71. The aspirated blood and fluid with the embolic debris from reservoir tip 15 can be passed through an extracorporeal filter and returned to a venous return cannula or arterial cannula 110.

The filter can also be deployed using obturator 75 as depicted in Fig. 8. Filter 10 is
20 mounted on a distal end of obturator 75. The obturator carrying the filter in a collapsed state is inserted into access lumen 66 to deploy filter 10 in aorta 101. Aspiration tube 20 extends proximally within obturator 75 and is connected through proximal end 76 of the obturator to pressure monitor 70 and vacuum 71. After completion of cardiovascular procedures, vacuum is disconnected proximally, and obturator 75 is withdrawn, removing
25 filter 10 and aspiration tube 20.

Fig. 9 depicts filter 10 mounted on distal end 79 of cannula 110. Aspiration tube 20 extends proximally within the cannula housing and is connected to pressure monitor 70 and vacuum 71. Alternatively, the aspiration tube may terminate just inside the vessel as shown also in Fig. 9. This embodiment applies as well to an aspiration tube carried by an
30 obturator or the handle of a filter. In this manner, air emboli, which tend to rise within the vessel, will migrate close to the aspiration port and be removed.

By way of example, when the filter disclosed herein is intended for use in the aorta, the area of the mesh required for the device is calculated from Bernoulli's equation as described in our earlier applications including Barbut et al., U.S. Application Serial No., U.S. Application Serial No. 08/553,137, filed November 7, 1995, Barbut et al., U.S. Application Serial No. 08/580,223, filed December 28, 1995, Barbut et al., U.S. Application Serial No. 08/584,759, filed January 9, 1996, Barbut et al., U.S. Patent No. 5,769,816, filed April 30, 1996, and Barbut et al., and U.S. Application Serial No. 08/645,762, filed May 14, 1996, all of which are incorporated herein by reference in their entirety.

10 In the embodiment of the filter that is to be used in the aorta, mesh with dimensions within the following ranges is desirable: mesh area is 0.5–10 in², more preferably 1–9 in², more preferably 2–8 in², more preferably 3–8 in², more preferably 4–8 in², more preferably 5–7 in²; mesh thickness is 60–280 μm, more preferably 70–270 μm, more preferably 80–260 μm, more preferably 90–250 μm, more preferably 100–250 μm, 15 more preferably 120–230 μm, more preferably 140–210 μm; thread diameter is 30–145 μm, more preferably 40–135 μm, more preferably 50–125 μm, more preferably 60–115 μm, more preferably 70–105 μm, and pore size is 500 μm or less, more preferably 400 μm or less, more preferably 300 μm or less, more preferably 200 μm or less, more preferably 100 μm or less, more preferably 50 μm or less and usually larger than at least a red blood 20 cell. In a preferred embodiment of the invention, mesh area is 2–8 in², mesh thickness is 60–200 μm, thread diameter is 30–100 μm, and pore size is 50–300 μm. In a further preferred embodiment of the invention, mesh area is 3–5 in², mesh thickness is 60–150 μm, thread diameter is 50–80 μm, and pore size is 100–250 μm.

During cardiovascular surgeries, it is known that distal embolization from embolic 25 debris generated from the proximal aorta occurs during removal of aortic cross-clamping. Occluder or dam 60 mounted on a distal end of insertion handle 65 can be utilized to minimize embolic complications as depicted in Fig. 10. In use, dam 60 is inserted into access lumen 66 and deployed in aorta 101 shortly before discontinuation of cardiopulmonary bypass. When aortic cross clamp 120 is removed, aspiration through 30 tube 20 is initiated and continued for approximately 30 seconds. In this way, blood and embolic material in the proximal aorta flow downstream into dam 60 and are aspirated and

removed, thereby preventing embolization to peripheral organs. After aspiration, dam 60 is collapsed and removed from aorta 101 to re-establish blood flow.

The length of the aspiration catheter will generally be between 10 and 100 centimeters for aortic use, preferably approximately between 20 and 50 centimeters. The inner diameter of the catheter will generally between 0.2 and 2 millimeters, preferably approximately between 0.5 and 1 millimeter. The filter will be capable of expanding to an outer diameter of at least 0.2 centimeters, more preferably at least 0.5 centimeters, more preferably at least 1.0 centimeters, more preferably at least 1.5 centimeters, more preferably at least 2.0 centimeters, more preferably at least 2.5 centimeters, more preferably at least 3.0 centimeters, more preferably at least 3.5 centimeters, more preferably at least 4.0 centimeters, more preferably at least 4.5 centimeters, more preferably at least 5.0 centimeters. The filter will be capable of contracting to an outer diameter of between 0.05 and 2.0 millimeters; preferably approximately between 0.8 and 1.2 millimeters. These ranges cover suitable diameters for both pediatric and adult use. The foregoing ranges are set forth solely for the purpose of illustrating typical device dimensions. The actual dimensions of a device constructed according to the principles of the present invention may obviously vary outside of the listed ranges without departing from those basic principles.

Although the foregoing invention has, for the purposes of clarity and understanding, been described in some detail by way of illustration and example, it will be obvious that certain changes and modifications may be practiced which will still fall within the scope of the appended claims. Moreover, it will be understood that each and every feature described for any given embodiment or in any reference incorporated herein, can be combined with any of the other embodiments described herein.

What is claimed is:

1. A filter, comprising:
a porous material arranged in a generally conical shape having an expandable base and a vertex;
5 a reservoir tip located at the vertex of the cone for collecting filtered embolic material; and
an aspiration tube having a proximal end, a distal end, and a lumen therebetween, the lumen communicating with a port at the distal end, the distal end of the aspiration tube extending into and communicating with the reservoir tip.
- 10 2. The filter of claim 1, wherein the reservoir tip comprises a nonporous material.
3. The filter of claim 1, further comprising a vacuum pump communicating with the proximal end of the aspiration tube.
4. The filter of claim 1, further comprising an infusion port in the reservoir tip
15 for infusion of saline.
5. The filter of claim 1, wherein the aspiration tube further comprises a turbine.
6. The filter of claim 2, wherein the reservoir tip comprises latex.
7. The filter of claim 6, wherein the latex covers a sector bounded by first and
20 second radii and the included arc of a circle.
8. The filter of claim 1, wherein the filter is mounted on a cannula.
9. The filter of claim 1, wherein the filter is mounted on an insertion handle.
10. The filter of claim 9, wherein the aspiration tube extends proximally within the insertion handle.

11. The filter of claim 1, wherein the aspiration tube extends proximally within an obturator.
12. The filter of claim 8, wherein the aspiration tube extends proximally within the cannula.
- 5 13. A method for protecting a patient from embolization, comprising the steps of:
- inserting a filter into the patient's aorta, the filter having a reservoir tip and an aspiration tube communicating with a port at the reservoir tip;
- expanding the filter;
- 10 aspirating emboli through the suction port, wherein embolic material is removed from the aorta; and
- collapsing the filter and removing the filter from the aorta.
14. The method of claim 13, further comprising the step of manipulating the aorta upstream of the filter.
- 15 15. The method of claim 13, further comprising the step of inserting an aortic cannula for perfusion of oxygenated blood.
16. The method of claim 15, further comprising the step of performing cardiopulmonary bypass.
17. The method of claim 13, further comprising the step of performing cardiac
- 20 surgery.
18. The method of claim 13, wherein the filter comprises a porous material arranged in a generally conical shape having an expanded base and a vertex.
19. The method of claim 13, wherein the reservoir tip collects filtered embolic material.
- 25 20. The method of claim 13, wherein the aspiration tube has a proximal end, a distal end, and a lumen therebetween.

21. The method of claim 13, wherein the port is at the distal end of the aspiration tube.

22. The method of claim 13, wherein the distal end of the aspiration tube extends into and communicates with the reservoir tip.

5 23. The method of claim 13, wherein the aspiration tube extends proximally within an obturator.

24. A vessel occluding device, comprising:
a nonporous material arranged in a generally conical shape having an expandable base and a vertex; and
10 an aspiration tube having a proximal end, a distal end, and a lumen therebetween, the lumen communicating with a port at the distal end, the distal end of the aspiration tube extending into and communicating with the vertex.

25. The device of claim 24, further comprising a reservoir tip located at the vertex of the cone for collecting filtered embolic material.

15 26. The device of claim 24, wherein the nonporous material is mounted at a distal end of a cannula.

27. The device of claim 24, wherein the nonporous material is mounted on an insertion handle.

28. The device of claim 24, wherein the aspiration tube is carried within the
20 cannula.

29. The device of claim 24, wherein the aspiration tube extends proximally within an obturator.

30. The device of claim 26, wherein the aspiration tube extends proximally within the cannula.

31. A method for protecting a patient from embolization, comprising the steps of:
- inserting a cannula into the patient's aorta, the cannula having a nonporous material and an aspiration tube communicating with a port at a vertex of the nonporous material;
- expanding the nonporous material to occlude the aorta;
- aspirating blood through the port, wherein embolic material is removed with the blood; and
- collapsing the nonporous material and removing the cannula from the aorta.
32. The method of claim 31, further comprising the step of manipulating the aorta upstream of the expanded nonporous material.
33. The method of claim 31, further comprising the step of inserting an aortic cannula for perfusion of oxygenated blood.
34. The method of claim 31, wherein the porous material is arranged in a generally conical shape having an expanded base and a vertex.
35. The method of claim 31, wherein a reservoir tip is located at the vertex of the cone for collecting filtered embolic material.
36. The method of claim 31, wherein the aspiration tube has a proximal end, a distal end, and a lumen therebetween.
37. The method of claim 35, wherein the aspiration tube extends into and communicates with the reservoir tip.
38. The method of claim 33, further comprising the step of performing cardiopulmonary bypass.
39. The method of claim 31, further comprising the step of performing cardiac surgery.
40. The method of claim 31, wherein the aspiration tube extends proximally within an obturator.

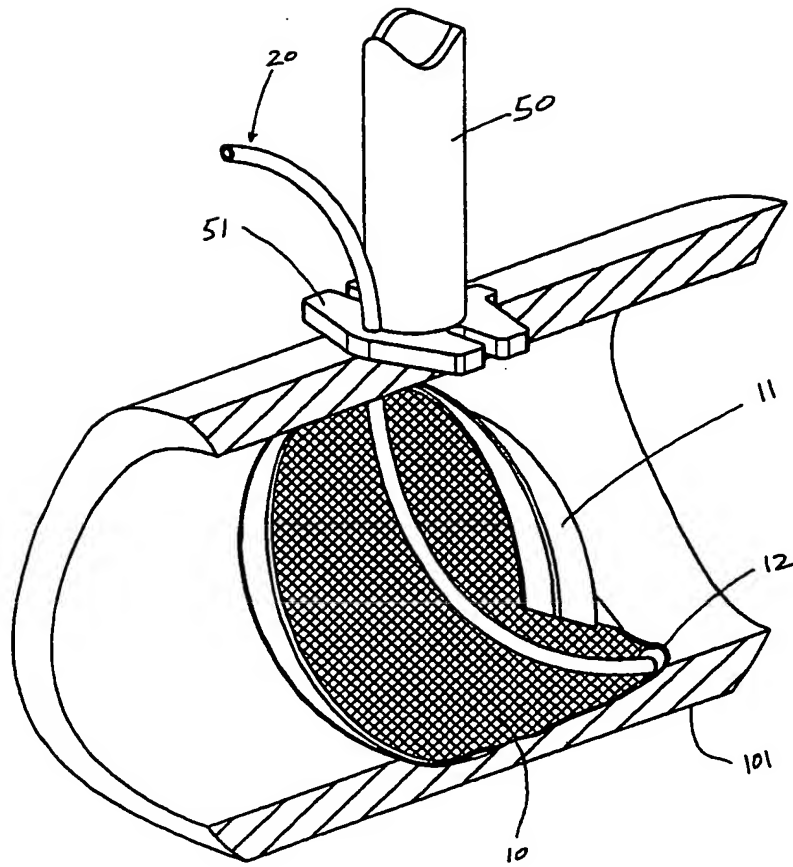


FIG. 1A

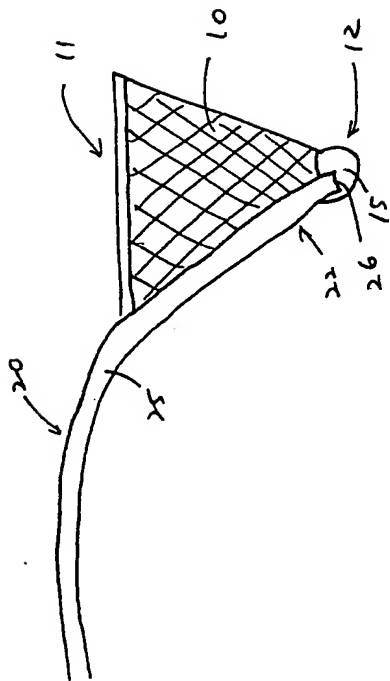


FIG. 1B

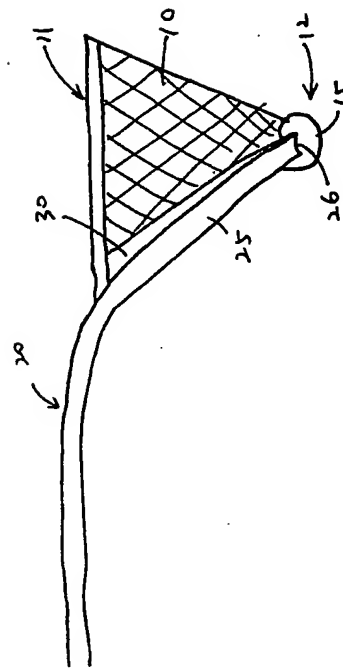


FIG. 2A

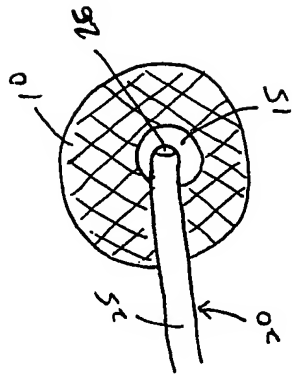


FIG. 1C

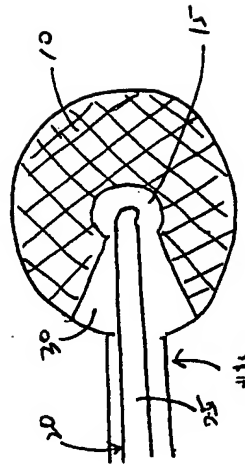


FIG. 2B

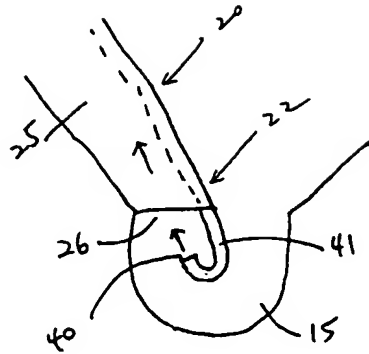


FIG. 3B

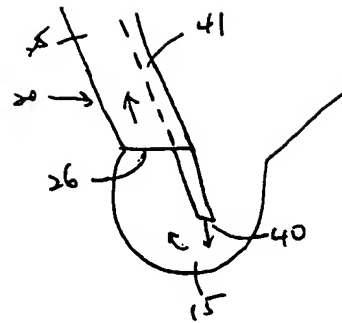


FIG. 3A

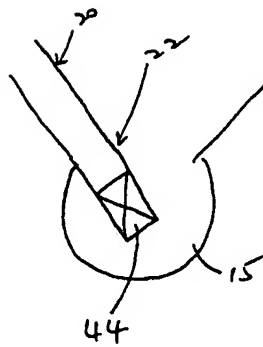


FIG. 4

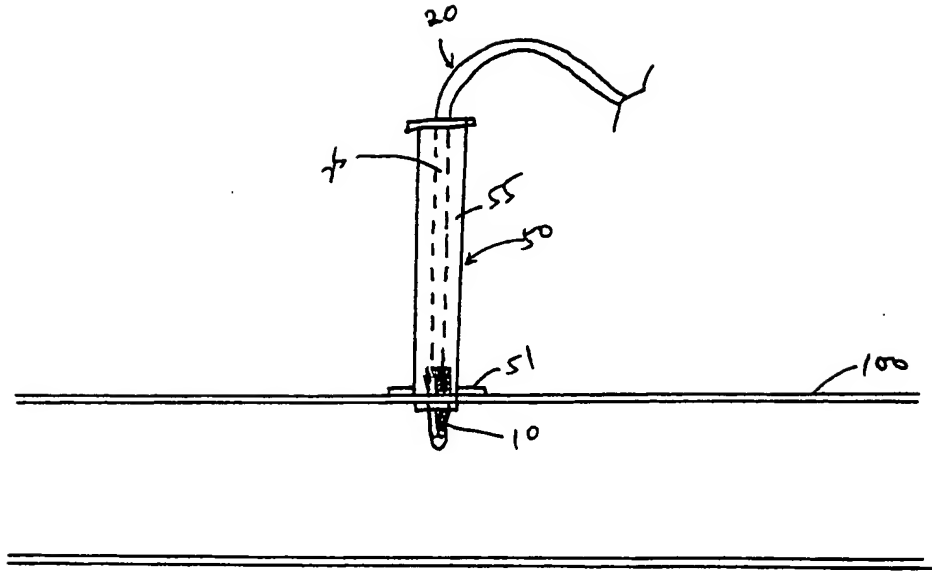


FIG. 5

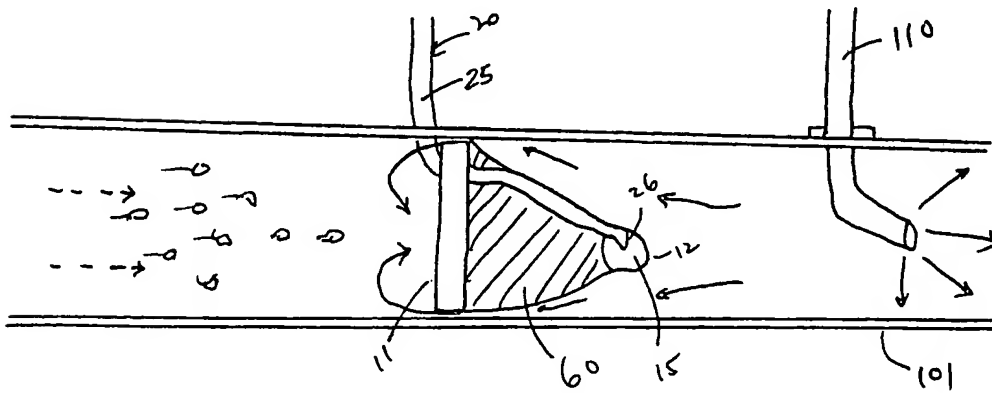


FIG. 6

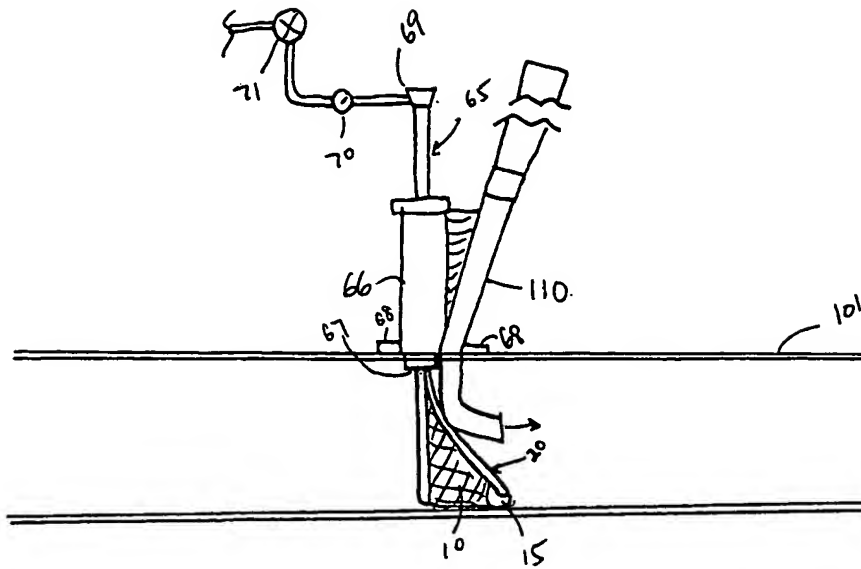


FIG. 7

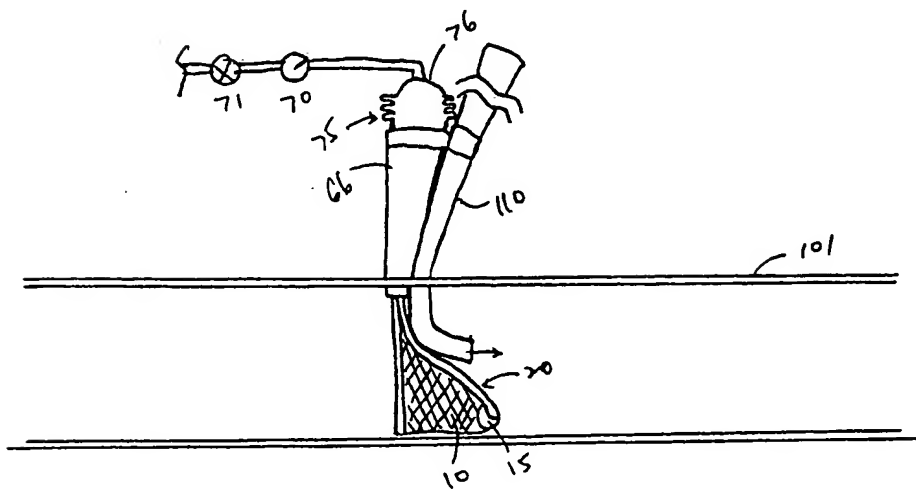


FIG. 8

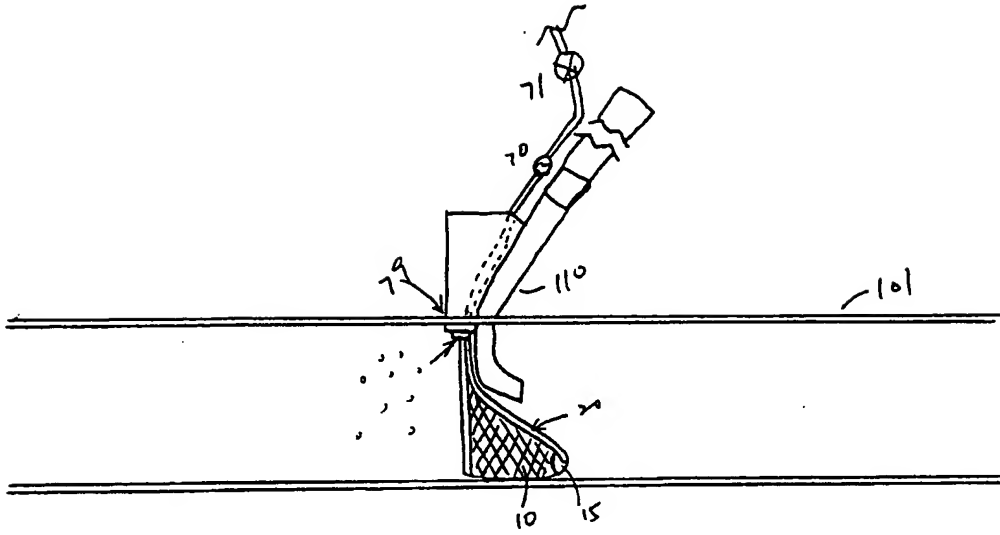


FIG. 9

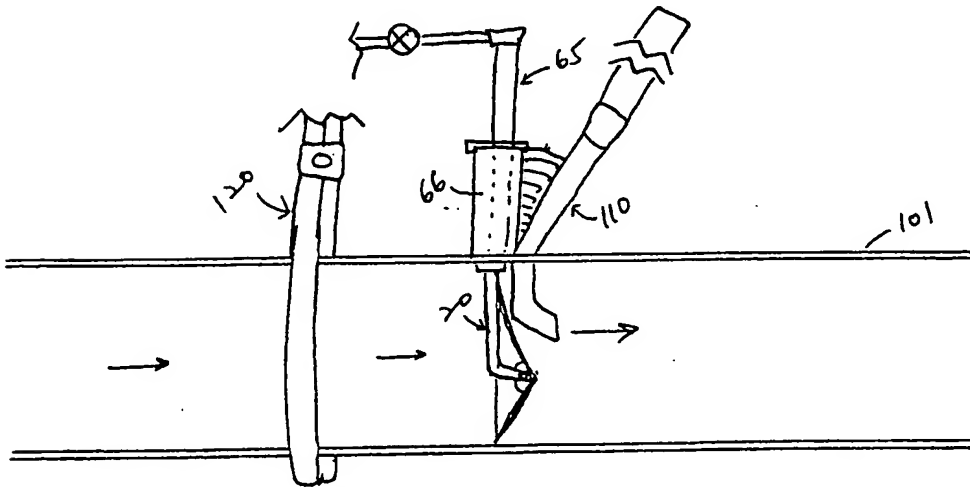


FIG. 10

INTERNATIONAL SEARCH REPORT

 International application No.
PCT/US00/35687

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) :A61M 1/00

US CL :604/540

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 604/540-541, 500, 507-510, 104

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X,P	US 6,136,016 A (BARBUT et al) 24 October 2000, see entire patent.	1-23
X	US 5,011,488 A (GINSBURG) 30 April 1991, see entire patent.	24-40
A,E	US 6,179,851 A (BARBUT et al) 30 January 2001, see entire patent.	1-40
A	US 5,865,802 A (YOON et al) 2 February 1999, see entire patent.	1-40
A	US 4,407,271 A (SCHIFF) 4 October 1983, see entire patent.	1-40
A	US 5,053,008 A (BAJAJ) 1 October 1991, see entire patent.	1-40

☒ Further documents are listed in the continuation of Box C.
 ☐ See patent family annex.

* Special categories of cited documents:	* T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
* A		document defining the general state of the art which is not considered to be of particular relevance
* E	* X	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
* L	* Y	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
* O		document referring to an oral disclosure, use, exhibition or other means
* P	* A	document published prior to the international filing date but later than the priority date claimed

Date of the actual completion of the international search

26 FEBRUARY 2001

Date of mailing of the international search report

27 APR 2001

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US00/35687

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4,723,549 A (WHOLEY et al) 9 February 1988, see entire patent.	1-40